

### **Support for Amendment**

New claims 54-64 are supported in part by the original claims and by the specification on page 9, line 26 through page 12, line 5; page 26, line 26 through page 27, line 17; page 49, line 4 through page 54, line 19; and page 57, line 29 through page 61, line 25. This amendment contains no new matter.

### **REMARKS**

Claims 1-53 are canceled without prejudice to their further prosecution in divisional and/or continuation applications. The claims now active in the application are specifically drawn to nucleic acid molecules that encode the polypeptide of SEQ ID NO:2. Claims 54-64 are active and pending in the application.

Sphingolipids serve as important signaling molecules to such diverse processes like cell growth, differentiation, and cell death. The novel sphingosine kinase (SphK) of the presently claimed invention contributes to signaling by forming sphingosine-1-phosphate, a potent inhibitor of cell death. Furthermore, over-expression of SphK inhibits chemotactic motility of several transformed cells line independently of cell surface receptors by acting through sphingosine-1-phosphate. Thus, SphK plays an important role in cellular regulation of processes linked to human disease, such as cancer. The claims of the present invention are drawn to a nucleic acid molecule that encodes a novel SphK polypeptide, which is represented by SEQ ID NO:2.

#### *Rejection of claims under 35 U.S.C. § 112, first paragraph*

The rejection of the claims as failing to satisfy the written description requirement and the enablement requirement under § 112, first paragraph has been obviated by appropriate amendment.

#### *Rejection of claims under 35 U.S.C. § 112, second paragraph*

The rejection of the claims as being indefinite under § 112, first paragraph has been obviated by amendment.

#### *Rejection of claims under 35 U.S.C. § 102*

The rejection of the claims as being anticipated by Young *et al.* and Kohama *et al.* under § 102, has been obviated by amendment.

*Objection of the claims*

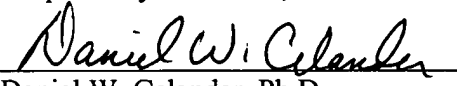
The objection of the claims based upon informalities has been obviated by amendment.

A copy of GenBank Acc. No. AA232646 that corresponds to reference C15 of the Information Disclosure Statement is provided. New corrected drawings of Figures 1-6 are also provided.

Applicants submit that the application is in condition for allowance. Early notice of such action is respectfully requested.

SONNENSCHN NATH & ROSENTHAL LLP  
P.O. Box #061080  
Wacker Drive Station  
Sears Tower  
Chicago, Illinois 60606-1080

Respectfully submitted,

  
Daniel W. Celander, Ph.D.  
Registration No. 52,710

Direct telephone calls to:

(312) 876-8071



Entrez

PubMed

Nucleotide

Protein

Genome

Structure

PMC

Taxonomy

Books

Search  for   

Limits

Preview/Index

History

Clipboard

Details

Show:

☐ 1: [AA232646](#). zr45f08.s1 Soares...[gi:1855648][Links](#)**IDENTIFIERS**

**dbEST Id:** 871993  
**EST name:** zr45f08.s1  
**GenBank Acc:** AA232646  
**GenBank gi:** 1855648  
**GDB Id:** 5428407

**CLONE INFO**

**Clone Id:** IMAGE:666375 (3')  
**Source:** IMAGE Consortium, LLNL  
**Insert length:** 1252  
**DNA type:** cDNA

**PRIMERS**

**Sequencing:** -41m13 fwd. ET from Amersham  
**PolyA Tail:** Unknown

**SEQUENCE**

```
CAGGCTGGGAATGTCACTTTATTTGGATTTGGTTTCGTGGGGTGGGGGTCTCAGAACAAAC
TAGAAGGCCTTACATAGGCAGCTGGGCCCAGCCAGCTGGCGTCTTGACCCAGGACTTCAT
TCTGGCCTGTCCCCCAAAGCATAGCCTCCACCTTCTCACCCTTCTCCAGAGGAGTCTCC
TCCACCCCCACAGGAGCTGTGGACAGGCCCTGCAGCCCTAGGGAAGGAGGAAGGGTCTTG
CAAGTAGACACTAAGGCACAGCGTGGGCCAGGGGTCTATAAGGGCTCTTCTGGCGGTGGCA
TCTGCTGGGGCTTCCAGCTGGGCGGGGGCTCCACGCAACCACTGACCATCCAGAAGTAGT
TTGGGTGCACCTGGCCCTGCACGGCCTCGCTAAC
```

**Entry Created:** Nov 27 1996  
**Last Updated:** Aug 6 1997

**COMMENTS**

This clone is available royalty-free through LLNL ; contact the IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.

**LIBRARY**

**Lib Name:** Soares\_NhHMPu\_S1  
**Organism:** Homo sapiens  
**Organ:** mixed (see below)  
**Tissue type:** Pooled human melanocyte, fetal heart, and pregnant uterus  
**Lab host:** DH10B  
**Vector:** pT7T3D-Pac (Pharmacia) with a modified polylinker  
**R. Site 1:** Not I  
**R. Site 2:** Eco RI  
**Description:** Equal amounts of plasmid DNA from three normalized libraries (melanocyte 2NbHM, pregnant uterus NbHPU, and fetal heart NbHH19W) were mixed, and ss circles were made in vitro.

Following HAP purification, this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from pools of 5,000 clones made from the same 3 libraries. The pools consisted of I.M.A.G.E. clones 260232-265223, 340488-345479, and 484488-489479.

**SUBMITTER**

Name: Wilson RK  
Institution: Washington University School of Medicine  
Address: 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
E-mail: [est@watson.wustl.edu](mailto:est@watson.wustl.edu)

**CITATIONS**

Title: WashU-Merck EST Project 1997  
Authors: Hillier,L., Allen,M., Bowles,L., Dubuque,T., Geisel,G., Jost,S., Kucaba,T., Lacy,M., Le,N., Lennon,G., Marra,M., Martin,J., Moore,B., Schellenberg,K., Steptoe,M., Tan,F., Theising,B., White,Y., Wylie,T., Waterston,R., Wilson,R.  
Year: 1997  
Status: Unpublished

**MAP DATA**

---

[Disclaimer](#) | [Write to the Help Desk](#)  
[NCBI](#) | [NLM](#) | [NIH](#)

Nov 18 2003 07:36:37